

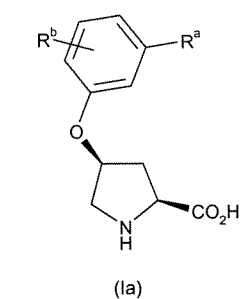
Listing of Claims

The following listing of Claims will replace all prior versions and listings of Claims in the Application.

Claims

1-10 (Cancelled)

11. (Currently Amended) A pharmaceutical composition comprising a compound of formula (1a) according to claim 14 :



wherein R^a is selected from halogen, hydroxy, (C_1-C_6) alkoxy, cyano, nitro, amino, hydroxycarbonyl, C_1-C_6 alkyl, C_1-C_6 alkenyl, C_1-C_6 alkynyl, hydroxy C_1-C_6 alkyl, C_1-C_6 alkoxy C_1-C_6 alkyl, perfluoro C_1-C_6 alkyl, perfluoro C_1-C_6 alkoxy, C_1-C_6 alkylamino, di- C_1-C_6 alkylamino, amino C_1-C_6 alkyl, C_1-C_6 alkylamino C_1-C_6 alkyl, di- C_1-C_6 alkylamino C_1-C_6 alkyl, C_1-C_6 acyl, C_1-C_6 acyloxy, C_1-C_6 acyloxy C_1-C_6 alkyl, C_1-C_6 acylamino, C_1-C_6 alkylthiocarbonyl, C_1-C_6 alkylthio, C_1-C_6 alkoxycarbonyl, C_1-C_6 alkylsulfonyl, C_1-C_6 alkylsulfonylamino, aminosulfonyl, C_1-C_6 alkylaminosulfonyl, di- C_1-C_6 alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R^b is selected from hydrogen, halogen, hydroxy, (C_1-C_6) alkoxy cyano, nitro, amino, hydroxycarbonyl, C_1-C_6 alkyl, C_1-C_6 alkenyl, C_1-C_6 alkynyl,

C₁-C₆ alkoxy, hydroxyC₁-C₆ alkyl, C₁-C₆ alkoxyC₁-C₆ alkyl, perfluoro C₁-C₆ alkyl, perfluoroC₁-C₆ alkoxy,

C₁-C₆ alkylamino, di- C₁-C₆ alkylamino, aminoC₁-C₆ alkyl, C₁-C₆ alkylaminoC₁-C₆ alkyl, di-C₁-C₆ alkylaminoC₁-C₆ alkyl,

C₁-C₆acyl, C₁-C₆acyloxy, C₁-C₆acyloxyC₁-C₆ alkyl, C₁-C₆ acylamino,

C₁-C₆ alkylthio, C₁-C₆ alkylthiocarbonyl, C₁-C₆ alkylthioxo, C₁-C₆ alkoxycarbonyl,

C₁-C₆ alkylsulfonyl, C₁-C₆ alkylsulfonylamino,

aminosulfonyl, C₁-C₆ alkylaminosulfonyl, di-C₁-C₆ alkylaminosulfonyl,

3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and

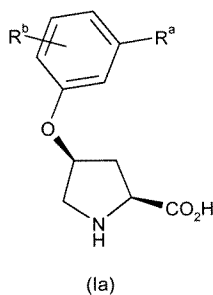
monocyclic heteroaryl;

or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

12. (Previously Presented) A combination comprising a compound of formula (Ia) according to claim 14, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

13. (Currently Amended) A combination according to claim 12, wherein the other therapeutically active agent is a PDEV inhibitor selected from sildenafil, vardenafil, tadalafil, 1-{6-ethoxy-5-[3-ethyl-6,7-dihydro-2-(2-methoxyethyl)-7-oxo-2H-pyrazolo[4,3- d]pyrimidin-5-yl]-3-pyridylsulfonyl}-4-ethylpiperazine, 5-(5-acetyl-2-butoxy-3-pyridinyl)-3-ethyl-2-(1-ethyl-3-azetidiny)-2,6-dihydro-7H-pyrazolo[4,3-*d*]pyrimidin-7-one and 5-[2-ethoxy-5-(4-ethylpiperazin-1-ylsulphonyl)pyridin-3-yl]-3-ethyl-2-[2-methoxyethyl]-2,6-dihydro-7H-pyrazolo[4,3-*d*]pyrimidin-7-one.

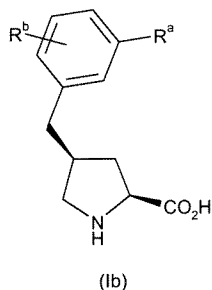
14. (Previously Presented) A compound of formula (Ia):



wherein R^a is selected from halogen, hydroxy, (C₁-C₆)alkoxy, cyano, nitro, amino, hydroxycarbonyl, C₁-C₆ alkyl, C₁-C₆ alkenyl, C₁-C₆ alkynyl, hydroxyC₁-C₆ alkyl, C₁-C₆ alkoxyC₁-C₆ alkyl, perfluoro C₁-C₆ alkyl, perfluoroC₁-C₆ alkoxy, C₁-C₆ alkylamino, di- C₁-C₆ alkylamino, aminoC₁-C₆ alkyl, C₁-C₆ alkylaminoC₁-C₆ alkyl, di-C₁-C₆ alkylaminoC₁-C₆ alkyl, C₁-C₆acyl, C₁-C₆acyloxy, C₁-C₆acyloxyC₁-C₆ alkyl, C₁-C₆ acylamino, C₁-C₆ alkylthiocarbonyl, C₁-C₆ alkylthioxo, C₁-C₆ alkoxycarbonyl, C₁-C₆ alkylsulfonyl, C₁-C₆ alkylsulfonylamino, aminosulfonyl, C₁-C₆ alkylaminosulfonyl, di-C₁-C₆ alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R^b is selected from hydrogen, halogen, hydroxy, (C₁-C₆)alkoxy cyano, nitro, amino, hydroxycarbonyl, C₁-C₆ alkyl, C₁-C₆ alkenyl, C₁-C₆ alkynyl, C₁-C₆ alkoxy, hydroxyC₁-C₆ alkyl, C₁-C₆ alkoxyC₁-C₆ alkyl, perfluoro C₁-C₆ alkyl, perfluoroC₁-C₆ alkoxy, C₁-C₆ alkylamino, di- C₁-C₆ alkylamino, aminoC₁-C₆ alkyl, C₁-C₆ alkylaminoC₁-C₆ alkyl, di-C₁-C₆ alkylaminoC₁-C₆ alkyl, C₁-C₆acyl, C₁-C₆acyloxy, C₁-C₆acyloxyC₁-C₆ alkyl, C₁-C₆ acylamino, C₁-C₆ alkylthio, C₁-C₆ alkylthiocarbonyl, C₁-C₆ alkylthioxo, C₁-C₆ alkoxycarbonyl, C₁-C₆ alkylsulfonyl, C₁-C₆ alkylsulfonylamino, aminosulfonyl, C₁-C₆ alkylaminosulfonyl, di-C₁-C₆ alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

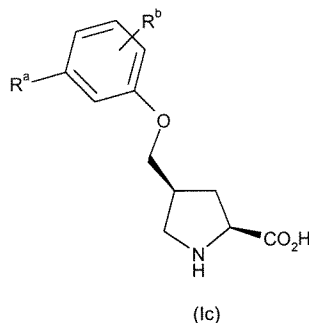
15. (Withdrawn) A compound of formula (Ib):



wherein R^a is selected from halogen, hydroxy, cyano, nitro, amino, hydroxycarbonyl, C₁-C₆ alkyl, C₁-C₆ alkenyl, C₁-C₆ alkynyl, hydroxyC₁-C₆ alkyl, C₁-C₆ alkoxyC₁-C₆ alkyl, perfluoroC₁-C₆ alkoxy, C₁-C₆ alkylamino, di-C₁-C₆ alkylamino, aminoC₁-C₆ alkyl, C₁-C₆ alkylaminoC₁-C₆ alkyl, di-C₁-C₆ alkylaminoC₁-C₆ alkyl, C₁-C₆acyl, C₁-C₆acyloxy, C₁-C₆acyloxyC₁-C₆ alkyl, C₁-C₆ acylamino, C₁-C₆ alkylthio, C₁-C₆ alkylthiocarbonyl, C₁-C₆ alkylthioxo, C₁-C₆ alkoxycarbonyl, C₁-C₆ alkylsulfonyl, C₁-C₆ alkylsulfonylamino, aminosulfonyl, C₁-C₆ alkylaminosulfonyl, di-C₁-C₆ alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R^b is selected from hydrogen, halogen, hydroxy, (C₁-C₆)alkoxy cyano, nitro, amino, hydroxycarbonyl, C₁-C₆ alkyl, C₁-C₆ alkenyl, C₁-C₆ alkynyl, C₁-C₆ alkoxy, hydroxyC₁-C₆ alkyl, C₁-C₆ alkoxyC₁-C₆ alkyl, perfluoro C₁-C₆ alkyl, perfluoroC₁-C₆ alkoxy, C₁-C₆ alkylamino, di-C₁-C₆ alkylamino, aminoC₁-C₆ alkyl, C₁-C₆ alkylaminoC₁-C₆ alkyl, di-C₁-C₆ alkylaminoC₁-C₆ alkyl, C₁-C₆acyl, C₁-C₆acyloxy, C₁-C₆acyloxyC₁-C₆ alkyl, C₁-C₆ acylamino, C₁-C₆ alkylthio, C₁-C₆ alkylthiocarbonyl, C₁-C₆ alkylthioxo, C₁-C₆ alkoxycarbonyl, C₁-C₆ alkylsulfonyl, C₁-C₆ alkylsulfonylamino, aminosulfonyl, C₁-C₆ alkylaminosulfonyl, di-C₁-C₆ alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

16. (Withdrawn) A compound of formula (Ic):



wherein R^a and R^b are independently selected from hydrogen, halogen, hydroxy, (C_1-C_6) alkoxy, cyano, nitro, amino, hydroxycarbonyl, C_1-C_6 alkyl, C_1-C_6 alkenyl, C_1-C_6 alkynyl, C_1-C_6 alkoxy, hydroxy C_1-C_6 alkyl, C_1-C_6 alkoxy C_1-C_6 alkyl, perfluoro C_1-C_6 alkyl, perfluoro C_1-C_6 alkoxy, C_1-C_6 alkylamino, di- C_1-C_6 alkylamino, amino C_1-C_6 alkyl, C_1-C_6 alkylamino C_1-C_6 alkyl, di- C_1-C_6 alkylamino C_1-C_6 alkyl, C_1-C_6 acyl, C_1-C_6 acyloxy, C_1-C_6 acyloxy C_1-C_6 alkyl, C_1-C_6 acylamino, C_1-C_6 alkylthio, C_1-C_6 alkylthiocarbonyl, C_1-C_6 alkylthioxo, C_1-C_6 alkoxycarbonyl, C_1-C_6 alkylsulfonyl, C_1-C_6 alkylsulfonylamino, aminosulfonyl, C_1-C_6 alkylaminosulfonyl, di- C_1-C_6 alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

17. (Withdrawn, Currently Amended) A compound of formula (1a) according to claim 4 14 which is:

(2S, 4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid;

Or a pharmaceutically acceptable salt thereof.

18. (Withdrawn) A compound of formula (1b) according to claim 15 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid;

(2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; and

(2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

19. (Withdrawn) A compound of formula (1c) according to claim 16 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-phenoxy)methyl-pyrrolidine-2-carboxylic acid;
(2S,4S)-4-(3,6-Difluoro-phenoxy)methyl-pyrrolidine-2-carboxylic acid;
(2S,4S)-4-(2,3-Difluoro-phenoxy)methyl-pyrrolidine-2-carboxylic acid; and
(2S,4S)-4-(3-Methoxy- phenoxy)methyl-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

20. (Withdrawn) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

21. (Withdrawn) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

22. (Withdrawn) A combination according to claim 21, wherein the other therapeutically active agent is a PDEV inhibitor.

23. (Withdrawn) A pharmaceutical composition comprising a compound of formula (1c) according to claim 16, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

24. (Withdrawn) A pharmaceutical composition comprising a compound of formula (1c) according to claim 16, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

25. (Withdrawn) A combination according to claim 24, wherein the other therapeutically active agent is a PDEV inhibitor.

26. (Withdrawn) The compound (2S, 4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid or a pharmaceutically acceptable salt thereof.